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LISTING OF CLAIMS

Claims 1-11 (CANCELED)

12- (NEW) A solid orodispersible pharmaceutical composition comprising:

- 5 - granules consisting of co-dried lactose and starch, and
 - piribedril or a pharmaceutically acceptable salt thereof.

13- (NEW) A composition according to claim 12, wherein the composition disintegrates in the mouth in less than three minutes.

14- (NEW) A composition according to claim 13, wherein the composition disintegrates in the mouth in less than one minute.

10 **15- (NEW)** A composition according to claim 12, comprising, in relation to the total weight of the composition :

- from 50 % to 95 % by weight of granules consisting of co-dried lactose and starch and
 - from 5 % to 50 % by weight of piribedil or a pharmaceutically acceptable salt thereof.

15 **16- (NEW)** A composition according to claim 15, comprising from 10 % to 20 % by weight of piribedil or a pharmaceutically acceptable salt thereof.

17- (NEW) A composition according to claim 12, further comprising one or more lubricants and a flow agent.

18- (NEW) A composition according to claim 12, further comprising citric acid.

20 **19- (NEW)** A composition according to claim 12, wherein the composition is in the form of a tablet.

20- (NEW) A tablet according to claim 19, wherein the tablet is obtained by direct compression.

21- (NEW) A tablet according to claim 20, wherein the tablet has a hardness from 15 to 50 Newtons.

22- (NEW) A tablet according to claim 21, wherein the tablet has a hardness of about 20 Newtons.

5 **23- (NEW)** A process for the manufacture of solid orodispersible compositions of piribedil, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than three minutes, wherein the piribedil, or a pharmaceutically acceptable salt thereof, is mixed with granules consisting of co-dried lactose and starch.

24- (NEW) A process for the manufacture of solid orodispersible compositions of piribedil, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than one minute, wherein the piribedil, or a pharmaceutically acceptable salt thereof, is mixed with granules consisting of co-dried lactose and starch.

10 **25- (NEW)** A method for treating a living animal body, including a human, afflicted with Parkinson's disease, including acute episodes of Parkinson's disease, comprising the step of administering to the living animal body, including a human, a composition according to claim 12, which is effective for treatment of Parkinson's disease, including acute episodes of Parkinson's disease.